Amendments to the Claims:

The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

1. (Currently amended) A method of treatment of a condition indicating treatment with a beta 4 subtype selective nicotinic acetylcholine receptor modulator comprising administering an effective amount of a compound represented by Formula (I) or pharmaceutically acceptable salts thereof:

$$\mathbb{R}^{4}$$
 $\mathbb{C}H_{2}$
 \mathbb{N}
 \mathbb{R}^{3}
 $\mathbb{C}H_{2}$
 \mathbb{N}
 \mathbb{R}^{1}

(I)

wherein:

$$R^1$$
 is -H,

 C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio, or

aryl-
$$C_{1-4}$$
alkyl;
 R^2 is -H,
-OH,
-C(O)-NH₂,
-NH₂,
-NH-Q-V-T, wherein Q is -C(O)-, -C(O)-NH-, -C(O)O-, or -SO₂-;

V is H, aryl, aryl- C_{1-12} alkyl, diaryl- C_{1-12} alkyl, lactonyl, or C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, - $C(O)OC_{1-4}$ alkyl, - $OC(O)C_{1-4}$ alkyl, aryl- C_{1-4} alkoxy, aryloxy, or SO_2C_{1-4} alkyl; and T is H, halogen, C_{1-5} alkyl, C_{1-4} alkoxy, nitro, aryl, aryl- C_{1-4} alkyl, or aryloxy unless V is H in which case T is absent; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia)

$$D \xrightarrow{E} Y \xrightarrow{Y} S \xrightarrow{(CH_2)_n} (CH_2)_n$$

$$W \xrightarrow{R^3} (Ia)$$

whereinD is O or S; and

E is O, S, NR⁵, C(R⁵)₂, O-CR⁵₂, NR⁵-CR⁵₂, NR⁵-CO, CR⁵₂-O, CR⁵₂-S(O)_r, CR⁵₂-NR⁵, CR⁵₂-CR⁵₂, CO-NR⁵, or CR⁵=CR⁵;

unless X is N in which case R2 is absent

R³ is

H, halogen, C_{1-4} alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF_3 , OC_{1-4} alkyl, aryl OC_{1-4} alkyl

carbonamido, -CO-, -CO₂H, -NH₂, NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋

4alkyl, or -C₁₋₄alkyl-OH;

R4 is

H, halogen, $C_{1.4}$ alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF_3 ,

OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy,

carbonamido, -CO-, -CO₂H, -NH₂, NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁.

4alkyl, or -C₁₋₄alkyl-OH;

R⁵ is each independently H or C_{1.4}alkyl;

X is

C or N;

W is C or N;

W' is C or N;

Y is C or N;

Y' is

C or N;

provided that there are no more than two N atoms in the aryl ring;

m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6;

provided that

when X, W, W', Y and Y' are all C, R^3 and R^4 are H and R^1 is selected from H, unsubstituted C_{1-4} alkyl and unsubstituted C_{3-4} cycloalkyl, R^2 may not be -OH;

and that

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when one of X, Y and Y' is N, R³ and R⁴ are H and R¹ is selected from H, unsubstituted C_{1.4}alkyl and unsubstituted C₃₋₄cycloalkyl, R² may not be H.

2. (Original) The method of claim 1 provided that when X, W, W', Y and Y' are all C and R³ and R⁴ are H, R² may not be -OH; and that when one of X, Y and Y' is N and R³ and R⁴ are H, R² may not be H.

3. (Currently amended) A method of treatment of dysfunctions of the central and autonomic nervous systems comprising administering an effective amount of a compound represented by Formula (I) or pharmaceutically acceptable salts thereof:

$$\mathbb{R}^{4}$$
 \mathbb{R}^{4}
 \mathbb{R}^{4}
 \mathbb{R}^{4}
 \mathbb{R}^{2}
 \mathbb{R}^{3}
 \mathbb{R}^{4}
 \mathbb{R}^{2}
 \mathbb{R}^{1}
 \mathbb{R}^{2}
 \mathbb{R}^{3}
 \mathbb{R}^{3}
 \mathbb{R}^{3}

(I)

wherein:

$$R^1$$
 is -H,

C₁₋₁₂alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C₁₋₄alkoxy or C₁₋₄alkylthio, or

V is H, aryl, aryl-C₁₋₁₂alkyl, diaryl-C₁₋₁₂alkyl, lactonyl, or C₁₋₁₈alkyl optionally substituted with halogen, $hydroxyl, C_{1-4}alkoxy, -C(O)OC_{1-4}alkyl, -OC(O)C_{1-4}alkyl, aryl-C_{1-4}alkoxy, aryloxy, or SO_2C_{1-4}alkyl; and all control of the c$ T is H, halogen, C₁₋₅alkyl, C₁₋₄alkoxy, nitro, aryl, aryl-C₁₋₄alkyl, or aryloxy unless V is H in which case T is absent; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia)

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whereinD is O or S; and

E is O, S, NR⁵, C(R⁵)₂, O-CR⁵₂, NR⁵-CR⁵₂, NR⁵-CO, CR⁵₂-O, CR⁵₂-S(O)₁, CR⁵₂-NR⁵, CR⁵₂-CR⁵₂, CO- NR^5 , or $CR^5=CR^5$;

unless X is N in which case R2 is absent

R³ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃,

OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy,

carbonamido, -CO-, -CO₂H, -NH₂, NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁-

4alkyl, or -C1-4alkyl-OH;

R4 is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃,

OC₁₄alkyl, aryloxy, arylC₁₄alkyl, arylC₁₄alkoxy, C₃,10</sub>cycloalkoxy, carboxy,

carbonamido, -CO-, -CO₂H, -NH₂, NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁.

4alkyl, or -C₁₋₄alkyl-OH;

R⁵ is each independently H or C₁₋₄alkyl;

X is C or N;

W is C or N;

W' is C or N;

Y is C or N;

Y' is C or N;

provided that there are no more than two N atoms in the aryl ring;

m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is $\frac{2}{3}$, $\frac{3}{4}$, $\frac{4}{5}$, or $\frac{6}{5}$;

provided that when X, W, W', Y and Y' are all C and R³ and R⁴ are H, R² may not be -OH; and that when one of X, Y and Y' is N and R³ and R⁴ are H, R² may not be H; and that when R² is H, OH or NH₂ and R³ and R⁴ are H, R¹ may not be aryl-C₁₋₄alkyl.

4. (Original) The method of any one of claims 1 to 3 wherein

 R^1 is -H, or C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio.

5. (Previously presented) The method of any one of claims 1 to 3, wherein

$$R^2$$
 is -H, -C(O)-NH₂,

-NH₂,

-NH-Q-V-T as defined in claim 1; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 1;

unless X is N in which case R2 is absent.

6. (Previously presented) The method of any one of claims 1 to 3, wherein R^2 is $-C(O)-NH_2$,

-NH-Q-V-T as defined in claim 1; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 1;

unless X is N in which case R² is absent.

7. (Previously presented) The method of any one of claims 1 to 3, wherein R^2 is $-C(O)-NH_2$,

V is as defined in claim 1; and

T is as defined in claim 1; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 1;

unless X is N in which case R2 is absent.

8-12. (Canceled)

13. A compound represented by Formula (I) or pharmaceutically acceptable salts thereof:

$$\mathbb{R}^{4}$$
 \mathbb{Y}
 \mathbb{Y}
 $\mathbb{C}H_{2}$
 \mathbb{N}
 \mathbb{R}^{3}
 $\mathbb{C}H_{2}$
 \mathbb{N}
 \mathbb{R}^{1}

(I)

wherein:

$$R^1$$
 is -H,

 C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio, or

V is H, aryl, aryl- C_{1-12} alkyl, diaryl- C_{1-12} alkyl, lactonyl, or C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, -C(O)OC₁₋₄alkyl, -OC(O)C₁₋₄alkyl, aryl- C_{1-4} alkoxy, aryloxy, or SO₂C₁₋₄alkyl; and T is H, halogen, C_{1-5} alkyl, C_{1-4} alkoxy, nitro, aryl, aryl- C_{1-4} alkyl, or aryloxy unless V is H in which case T is absent; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia)

whereinD is O or S; and

E is O, S, NR⁵, C(R⁵)₂, O-CR⁵₂, NR⁵-CR⁵₂, NR⁵-CO, CR⁵₂-O, CR⁵₂-S(O)_r, CR⁵₂-NR⁵, CR⁵₂-CR⁵₂, CO-NR⁵, or CR⁵=CR⁵;

unless X is N in which case R² is absent

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R³ is H, halogen, C_{1-4} alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF_3 , OC_{1-4} alkyl, aryloxy, aryl C_{1-4} alkyl, aryl C_{1-4} alkoxy, C_{3-10} cycloalkoxy, carboxy, carbonamido, -CO-, -CO₂H, -NH₂, NH- C_{1-4} alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁. 4alkyl, or - C_{1-4} alkyl-OH;

R⁴ is H, halogen, C_{1-4} alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF_3 , OC_{1-4} alkyl, aryloxy, aryl C_{1-4} alkyl, aryl C_{1-4} alkoxy, C_{3-10} cycloalkoxy, carboxy, carbonamido, -CO-, -CO₂H, -NH₂, NH- C_{1-4} alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁. 4alkyl, or - C_{1-4} alkyl-OH;

R⁵ is each independently H or C₁₋₄alkyl;

X is C or N;

W is C or N;

W' is C or N;

Y is C or N;

Y' is C or N;

provided that there are no more than two N atoms in the aryl ring;

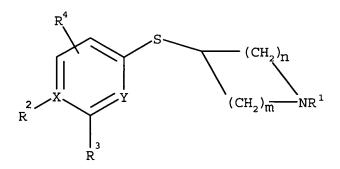
m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6;

provided that when X, W, W', Y and Y' are all C and R³ and R⁴ are H, R² may not be -OH; and that

when one of X, Y and Y' is N and R^3 and R^4 are H, R^2 may not be H; and that when R^2 is H, OH or NH₂ and R^3 and R^4 are H, R^1 may not be aryl-C₁₋₄alkyl; and excluding compounds represented by Formula I'' or pharmaceutically acceptable salts thereof:



(I")

wherein:

 R^1 , X, Y, m and n are as defined above R^2 is -H,

 $-NH_2$,

-NH-Q-V-T, wherein Q is -C(O)- or -SO₂- and

V and T are as defined above;

unless X is N in which case R2 is absent

R³ is H, halogen, C₁₋₄alkyl, OC₁₋₄alkyl, -NH₂, NH-C₁₋₄alkyl, or hydroxy;

R⁴ is H, halogen, C₁₋₄alkyl, OC₁₋₄alkyl, CO₂H, -NH₂, NH-C₁₋₄alkyl, or hydroxy.

14. (Original) A compound as claimed in claim 13 wherein

R¹ is -H, or

 $C_{1\text{-}12} alkyl \ optionally \ substituted \ with \ 1, 2 \ or 3 \ groups \ independently \ selected \ from halogen, hydroxyl, thiol, $C_{1\text{-}4} alkoxy \ or \ C_{1\text{-}4} alkylthio.$

15. (Original) A compound as claimed in claim 13 or claim 14, wherein

 R^2 is -H,

-C(O)-NH₂,

-NH₂,

-NH-Q-V-T as defined in claim 13; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 13;

unless X is N in which case R2 is absent.

16. (Original) A compound as claimed in any one of claims 13 to 15, wherein

 R^2 is $-C(O)-NH_2$,

-NH-Q-V-T as defined in claim 13; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 13:

unless X is N in which case R2 is absent.

17. (Previously presented) A compound as claimed in any one of claims 13 to 14,

wherein

R² is -C(O)-NH₂, -NH-Q-V-T, whereinQ is -C(O)-NH-, or -C(O)O-;

V is as defined in claim 13; and

T is as defined in claim 13; or

linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia) as defined in claim 13;

unless X is N in which case R² is absent.

18. (Currently amended) A compound as claimed in claim 13 which is represented by Formula (II) or pharmaceutically acceptable salts thereof:

$$\mathbb{R}^{4}$$

$$\mathbb{C}H_{2})_{n}$$

$$\mathbb{R}^{2}$$

$$\mathbb{R}^{3}$$

$$\mathbb{C}H_{2})_{m}$$

$$\mathbb{N}\mathbb{R}^{1}$$

(II)

wherein:

R1 is -H; or

 C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio; or aryl- C_{1-4} alkyl;

aryı-C₁.

 R^2 is -H;

-OH;

 $-C(O)-NH_2$

-NH₂;

-NH-Q-V-T

Q is -C(O)-;

-C(O)-NH-;

-C(O)O-; or

-SO₂-

V is aryl;

aryl- C_{1-12} alkyl;

diaryl-C₁₋₁₂ alkyl;

lactonyl; or

 $C_{1\text{--}18} \text{ alkyl optionally substituted with halogen, hydroxyl, } C_{1\text{--}4} \text{ alkoxy, -C(O)OC}_{1\text{--}4} \text{ alkyl, -OC(O)C}_{1\text{--}4} \text{ alkyl, aryl-C}_{1\text{--}4} \text{ alkoxy, aryloxy, } SO_2C_{1\text{--}4} \text{ alkyl;}$

T is H;

halogen;

aryl;

aryl- C_{1-4} alkyl; or aryloxy;

unless X is N in which case R2 is absent

 R^3 and R^4 are each independently selected from H, halogen, C_{1-4} alkyl, cyano, CF_3 , OC_{1-4} alkyl, aryloxy, aryl C_{1-4} alkoxy, C_{3-10} cycloalkoxy, carboxy, carbonamido, -CO-, -CO₂H, -NH₂, NH- C_{1-4} alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄ alkyl, -C₁₋₄ alkyl-OH; X is C or N;

W is C or N, provided that both X and Y are not N;

Y is C or N

m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6.

- 19. (Original) A compound as claimed in claim 18 wherein R^1 is H; C_{1-6} alkyl optionally substituted with 1 or 2 hydroxyl groups; or aryl- C_{1-4} alkyl.
- 20. (Original) A compound as claimed in claim 19 wherein R¹ is benzyl, p-methoxybenzyl, furanylmethyl, imidazolylmethyl, pyridinylmethyl, thienylmethyl, pyridylmethyl, N-hydroxypyridylmethyl or thiazolylmethyl.
- 21. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is H, R³ is carbonamido (-CONH₂) or C₁₋₄ alkyl-OH, and R⁴ is H, C₁₋₄alkyl, CF₃, halogen or cyano.
- 22. (Original) A compound as claimed in any one of claims 18 to 20 wherein R^2 is OH, and R^3 and R^4 each independently represent H, C_{1-4} alkyl, CF_{3} , cyano or halogen.
- 23. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-Q-V-T; T is H and R³ and R⁴ each independently represent H, methyl, CF₃, chloro- or cyano-.
- 24. (Original) A compound as claimed in any one of claims 18 to 20 wherein R^2 is of formula –NH-SO₂-V-T; V is aryl, -C₁₋₁₂ alkyl or aryl-C₁₋₁₂ alkyl; R^3 is H, methyl, CF₃, Cl or cyano and R^4 is H.

25. (Original) A compound as claimed in any one of claims 18 to 20 wherein R^2 is of formula –NH-SO₂-V-T, V is selected from C_{1-12} alkyl, phenyl, naphthyl, thienyl, oxazolyl, isoxazolyl, or phenyl(CH=CH)–, optionally substituted with 1, 2, 3 or 4 substituents selected from:

```
-NO<sub>2</sub>;
halogen;
-CF<sub>3</sub>;
C<sub>1-12</sub> alkoxy;
C<sub>1-12</sub> alkylthio;
C<sub>1-12</sub> alkyl;
C<sub>1-4</sub> alkylsulfonyl;
-CN;
-OCF<sub>3</sub>;
-C(O)OC<sub>1-4</sub> alkyl;
-OCH<sub>2</sub>CF<sub>3</sub>;
-NHC(O) C<sub>1-4</sub> alkyl.
```

26. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-SO₂-V-T, T is selected from H; or diazole, oxazole, isoxazole, phenyl or phenoxy, optionally substituted with 1, 2, 3 or 4 substituents selected from

-NO₂; halogen; -CF₃; C₁₋₁₂ alkoxy; C₁₋₁₂ alkylthio; C₁₋₁₂ alkyl; C₁₋₄ alkylsulfonyl; -CN; -OCF₃; -C(O)OC₁₋₄ alkyl; -OCH₂CF₃; -NHC(O) C₁₋₄ alkyl.

27. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-SO₂-V-T, V is selected from 3-chloro-4-methylphenyl, 3-chlorophenyl, 3-methoxyphenyl, 4-bromophenyl, 4-methoxyphenyl, 4-methylphenyl, naphthyl, 2,4,6-trimethylphenyl, phenyl(CH=CH)-,

4-chlorophenyl, 2-chlorophenyl, 2,5-dichlorothien-3-yl, 2,5,6-trimethyl-4-methoxyphenyl, 4-methoxyphenyl, 2,3,4-trifluorophenyl, 3-cyanophenyl, 2-methoxycarbonylthien-3-yl or 4-pentylphenyl and T is H.

28. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-SO₂-V-T, T is 2-chloro-5-nitrophenoxy and V is phenyl.

29. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-C(O)-V-T wherein V is selected from

aryl; aryl- C_{1-12} alkyl; diaryl- C_{1-12} alkyl; lactonyl; or C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, $C(O)OC_{1-4}$ alkyl, $OC(O)C_{1-4}$ alkyl, aryl- C_{1-4} alkoxy, aryloxy.

30. (Original) A compound as claimed in any one of claims 18 to 20 wherein R^2 is of formula –NH-C(O)-V-T, and V is selected from C_{1-12} alkyl, phenyl, phenyl- C_{1-12} alkyl, diphenylmethyl, naphthyl, furanyl, thienyl, diazolyl, pyridinyl, thiazolyl, benzothienyl, fluorenyl, oxazolyl or isoxazolyl, optionally substituted with 1, 2, 3 or 4 substituents independently selected from

-NO₂; halogen; -CF₃; C₁₋₁₂ alkoxy; C₁₋₁₂ alkylthio; C₁₋₁₂ alkyl; C₁₋₄ alkylsulfonyl; -CN; -OCF₃; -C(O)O-C₁₋₄ alkyl; -OCH₂CF₃.

31. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-C(O)-V-T, T is selected from

н;

halogen; or

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with 1, 2, 3 or 4 substituents selected from -NO₂; halogen; -CF₃; C₁₋₁₂ alkylthio; C₁₋₁₂ alkoxy; C_{1-12} alkyl; C₁₋₄ alkylsulfonyl; -CN; -OCF₃; -C(O)O-C₁₋₄ alkyl. 32. (Original) A compound as claimed in any one of Claims 18 to 20 wherein R² is of formula -NH-C(O)N-V-T wherein V is selected from C₁₋₁₈ alkyl optionally substituted with halogen, hydroxyl, C₁₋₄ alkoxy, C(O)OC₁₋₄ alkyl, OC(O)C₁₋₄ alkyl, aryl-C₁.alkoxy, aryloxy; aryl; or aryl-C₁₋₁₂ alkyl. 33. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of formula –NH-C(O)NH-V-T, V is selected from phenyl, phenyl-C₁₋₁₂ alkyl or naphthyl optionally substituted with 1, 2, 3 or 4 substituents selected from $-NO_2$; halogen; $-CF_3$; C₁₋₁₂ alkylthio; C_{1-12} alkoxy; C_{1-12} alkyl; C₁₋₄ alkylsulfonyl; -CN; -OCF₃; -C(O)O-C₁₋₄ alkyl. 34. (Original) A compound as claimed in any one of claims 18 to 20 wherein R² is of

formula -NH-C(O)O-V-T, wherein V is selected from

diazole, oxazole, isoxazole, phenyl, phenoxy or benzodioxanyl optionally substituted

 C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, $C(O)OC_{1-4}$ alkyl, $OC(O)C_{1-4}$ alkyl, aryl- C_{1-4} alkoxy, aryloxy; aryl; or aryl- C_{1-12} alkyl.

35. (Original) A compound as claimed in any one of claims 18 to 20 wherein R^2 is of formula –NH-C(O)O-V-T, preferably V is selected from phenyl or phenyl- C_{1-12} alkyl optionally substituted with 1, 2, 3 or 4 substituents selected from

 $-NO_2$;

halogen;

 $-CF_3$;

C₁₋₁₂ alkylthio;

 C_{1-12} alkoxy;

C₁₋₁₂ alkyl;

C₁₋₄ alkylsulfonyl;

-CN;

-OCF₃;

-C(O)O-C₁₋₄ alkyl; or

-OCH₂CF₃.

36. (Original) A compound as claimed in claim 13 wherein R² is of formula –NH-C(O)-

V-T

wherein

V is H, C₁₋₆alkyl, C₃₋₆cycloalkyl, aryl or aryl-C₁₋₁₂alkyl; and

T is H, halogen, C_{1.5}alkyl, C_{1.4}alkoxy, nitro, aryl, aryl-C_{1.4}alkyl, or aryloxy unless V is H in which case T is absent.

37. (Original) A compound as claimed in claim 36

wherein

V is H, C₁₋₆alkyl or C₃₋₆cycloalkyl, and

T is H unless V is H in which case T is absent.

38. (Orignal) A compound as claimed in claim 36

wherein

V is aryl or aryl-C₁₋₁₂alkyl, and

T is H, halogen, C₁₋₅alkyl, C₁₋₄alkoxy, nitro, aryl, aryl-C₁₋₄alkyl, or aryloxy.

39. (Original) A compound as claimed in claim 38

wherein

V is phenyl, pyridyl, thienyl, thiazolyl, thiadiazolyl, or phenyl-C₁₋₆alkyl; and

T is H, halogen, C₁₋₅alkyl, C₁₋₄alkoxy, nitro, aryl, aryl-C₁₋₄alkyl, or aryloxy.

40. (Currently amended) A compound as claimed in claim 13

wherein

R¹ is -H, $C_{1-12} \text{alkyl optionally substituted with 1, 2 or 3 groups independently selected} \\ \text{from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio, or <math display="block"> \text{aryl-C_{1-4} alkyl;}$

R² is -NH₂, or -NH-Q-V-T, wherein Q is -C(O)-, -C(O)-NH-, -C(O)O-, or -SO₂-; V is H, aryl, aryl- C_{1-12} alkyl, diaryl- C_{1-12} alkyl, lactonyl, or C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, -C(O)OC₁₋₄alkyl, -OC(O)C₁. 4alkyl, aryl- C_{1-4} alkoxy, aryloxy, or SO₂C₁₋₄alkyl; and T is H, halogen, aryl, aryl- C_{1-4} alkyl, or aryloxy unless V is H in which case T is

absent,

- R³ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄alkyl, or -C₁₋₄alkyl-OH;
- R⁴ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄alkyl, or -C₁₋₄alkyl-OH;

X is C;

W is C or N;

W' is C or N;

Y is C or N;

Y' is C or N;

provided that there are not more than two N atoms in the aryl ring and provided that at least one of W, W', Y or Y' is N;

m is 1, 2, or 3; n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6.

41. (Original) A compound as claimed in claim 40

wherein

W is C;

W' is C;

Y' is C; and

Y is N.

42. (Original) A compound as claimed in claim 40

wherein

W is N;

W' is C;

Y' is C; and

Y is C.

43. (Original) A compound as claimed in any one of claims 40 to 42 wherein R^2 is -NH₂.

44. (Original) A compound as claimed in any one of claims 40 to 42

wherein

 R^2 is -NH-Q-V-T, wherein Q is -C(O)-, -C(O)-NH-, -C(O)O-, or -SO₂-;

V is H, aryl, aryl- C_{1-12} alkyl, diaryl- C_{1-12} alkyl, lactonyl, or C_{1-18} alkyl optionally substituted with halogen, hydroxyl, C_{1-4} alkoxy, - $C(O)OC_{1-4}$ alkyl, - $OC(O)C_{1-4}$ alkyl, aryl- C_{1-4} alkoxy, aryloxy, or SO_2C_{1-4} alkyl; and T is H, halogen, aryl, aryl- C_{1-4} alkyl, or aryloxy unless V is H in which case T is absent.

45. (Original) A compound as claimed in claim 44 wherein Q is -SO₂- or -CO-.

46. (Currently amended) A compound as claimed in Claim 13

wherein:

 R^1 is -H,

 C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} alkoxy or C_{1-4} alkylthio, or aryl- C_{1-4} alkyl;

R² is linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia)

(la)

whereinD is O or S; and

E is O, S, NR⁵, or $C(R^5)_2$,

R³ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄alkyl, or -C₁₋₄alkyl-OH;

R⁴ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄alkyl, or -C₁₋₄alkyl-OH;

R⁵ is each independently H or C₁₋₄alkyl;

X is

C;

W is C or N;

W' is C;

Y is C or N;

Y' is C or N;

provided that there are no more than two N atoms in the aryl ring,

m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6.

47. A compound as claimed in Claim 46 wherein E is O or NR⁵.

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48. A compound as claimed in Claim 46 or 47 wherein R^5 is/are each independently H or C_{1-4} alkyl.

49. A compound as claimed in Claim 13

wherein:

R¹ is -H.

 C_{1-12} alkyl optionally substituted with 1, 2 or 3 groups independently selected from halogen, hydroxyl, thiol, C_{1-4} al

koxy or C₁₋₄alkylthio, or aryl-C₁₋₄alkyl;

R² is linked back to the aromatic ring so as to form a fused bicyclic compound represented by Formula (Ia)

whereinD is O or S; and

 $E \text{ is O-CR}^{5}_{2}, NR^{5}-CR^{5}_{2}, NR^{5}-CO, CR^{5}_{2}-O, CR^{5}_{2}-S(O)_{r}, CR^{5}_{2}-NR^{5}, CR^{5}_{2}-CR^{5}_{2}, CO-NR^{5}, or CR^{5}=CR^{5};$

R³ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁.

4alkyl, or -C₁₋₄alkyl-OH;

R⁴ is H, halogen, C₁₋₄alkyl optionally substituted with from 1 to 3 fluorine atoms, cyano, CF₃, OC₁₋₄alkyl, aryloxy, arylC₁₋₄alkyl, arylC₁₋₄alkoxy, C₃₋₁₀cycloalkoxy, carboxy, carbonamido, -CO-NH-C₁₋₄alkyl, aryl, hydroxy, -SO₂NH₂, -SO₂NHC₁₋₄alkyl, or -C₁₋₄alkyl-OH;

R⁵ is each independently H, C₁₋₄alkyl;

X is C;

W is C or N;

W' is C;

Y is C or N;

Y' is C or N;

provided that there are no more than two N atoms in the aryl ring;

m is 1, 2, or 3;

n is 1, 2, or 3; and

the sum of m and n is 2, 3, 4, 5, or 6.

- 50. A compound as claimed in Claim 49 wherein E is O-CR⁵₂, NR⁵-CR⁵₂, NR⁵-CO, CR⁵₂-CR⁵₂, or CR⁵=CR⁵.
- 51. A compound as claimed in Claim 49 or 50 wherein E is O-CR⁵₂, NR⁵-CO, or CR⁵=CR⁵.
- 52. (previously presented) A compound as claimed in any one of Claims 49 to 50 wherein R^5 is/are each independently H or C_{1-4} alkyl.
 - 53. (canceled)
- 54. (previously presented) A compound as claimed in any one of claims 18 to 20 wherein m is 2 and n is 2.
- 55. (previously presented) A compound as claimed in any one of claims 18 to 20 wherein X, Y and W are C.
 - 56. (canceled)
- 57. (previously presented) A pharmaceutical composition comprising a compound of claim 13 with a pharmaceutically acceptable diluent or carrier.